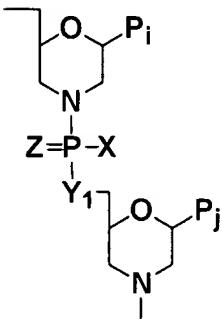


**Amendments to the Claims:**

1-27. (Cancelled)

28. (Currently amended) A method for treating a vascular injury site in a human patient by reducing restenosis at the site, said method comprising:

administering to the patient, by intravascular delivery directly to the vascular injury site, a morpholino antisense compound having uncharged phosphorodiamidate intersubunit linkages of the form:



where  $X=N(CH_3)_2$ ,  $Y=O$ , and  $Z=O[;]$ , and  $P_i$  and  $P_j$  are independently selected from adenine, guanine, cytosine, thymine and uracil; and comprising the sequence identified as SEQ ID NO: 1, in an amount effective to reduce restenosis in the patient.

29-31. (Cancelled)

32. (Previously presented) The method of claim 28, wherein said administering is carried out by injecting the antisense compound from an injection balloon catheter directly into the vascular injury site, under pressure, through injectors contained on the surface of the catheter balloon, wherein the vascular injury site comprises a vascular wall having a tunica media and wherein said injectors are capable of penetrating the tunica media in the vascular wall.

33. (Previously presented) The method of claim 32, wherein the catheter balloon has a plurality of outer-facing channels that are connected to a drug-delivery lumen of the catheter, each channel having one or more injection ports, and said injecting includes forcing

a solution or suspension of the antisense compound from said drug-delivery lumen through said injection ports when the balloon is in an inflated position.

34. (Previously presented) The method of claim 33, wherein the amount of antisense compound administered is between 5 and 20 mg.

35. (Previously presented) The method of claim 28, wherein said administering is carried out by contacting the vascular injury site with an intravascular stent having a coating containing the antisense compound in diffusible form.

36. (Previously presented) The method of claim 35, wherein the coating is designed to release the majority of the antisense compound in the coating over a period of 5-60 minutes following balloon angioplasty.

37. (Previously presented) The method of claim 36, wherein the intravascular stent is biodegradable.

38. (Previously presented) The method of claim 28, wherein said administering comprises placing the antisense compound in direct contact with the region of injury, in an amount effective to deliver between about 0.5 and 2 mg antisense compound into the vessel tissue.

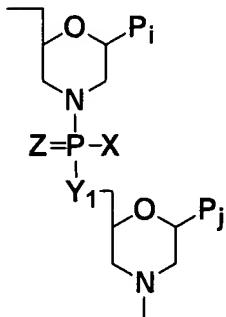
39. (Previously presented) The method of claim 38, wherein the compound is derivatized with a moiety that enhances the solubility of the antisense compound in aqueous medium, and the antisense compound is administered from a solution containing at least about 30 mg/ml of the antisense compound.

40. (Previously presented) The method of claim 39, wherein said moiety is triethyleneglycol attached to the 5' end of the antisense compound.

41. (Currently amended) A method for treating a vascular injury site in a human patient, said method comprising:

providing an intravascular stent wherein said stent or a coating on said stent contains a

morpholino antisense compound in diffusible form, wherein the morpholino antisense compound has uncharged phosphorodiamide intersubunit linkages of the form:



where  $X=N(CH_3)_2$ ,  $Y=O$ , and  $Z=O$ ,  $[[;]]$ , and  $Pi$  and  $Pj$  are independently selected from adenine, guanine, cytosine, thymine and uracil; and comprises the sequence identified as SEQ ID NO: 1, and

contacting the vascular injury site with said stent, effective to administer said compound to the patient, in an amount effective to reduce restenosis in the patient.

42-44. (Cancelled)

45. (Previously presented) The method of claim 41, wherein the coating is designed to release the majority of the antisense compound in the coating over a period of 5-60 minutes following balloon angioplasty.

46. (Previously presented) The method of claim 41, wherein the stent is biodegradable.

47. (Previously presented) The method of claim 41, wherein the compound is derivatized with a moiety that enhances the solubility of the compound in aqueous medium, to a level of at least about 30 mg/ml of the antisense compound.

48. (Previously presented) The method of claim 47, wherein said moiety is triethyleneglycol attached to the 5' end of the compound.

49. (Previously presented) The method of claim 41, wherein said contacting comprises placing the stent at the vessel site at the time of balloon angioplasty or during coronary bypass surgery.